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- (54) Silicone derivatives, their production and use
- (57) Disclosed are amino silicone derivatives having triazinyl groups or pyrimidinyl groups bound to amino groups of amino silicones, their production and use, said silicone derivatives being able to gel silicone oil or to increase the viscosity thereof stably and homogeneously, and therefore being useful as bases for cosmetics, pharmaceutical preparations and industrial materials.

#### Description

#### FIELD OF THE INVENTION

The present invention relates to novel silicone derivatives, their production and use.

# BACKGROUND OF THE INVENTION

Slicone oil is useful for its heat resistance, lubrishy, water repellency, gloss retention, miss prevention, artistato properities, most release lubricant, comorsion resistance, nethmical sability and safety. Utilism prince properities, most oil has hitherto been used as bases of compositions in a wide variety of fileds including various industrial materials such as textile heading agents, sursion obthicratiss, water repelling agents, resim confiders, patin address, electrical relating agents, heat media, grease, oil for machinery, toam stabilizers and anti-loam agents, pharmaceutical preparations and cosmelos.

In particular, allicone oil has been used as bases of pharmaceuscial preparations and cournelies because of its safety. Usually, low-viscosity sitions oil having a viscosity of 100 cor o less at mon temperature is widely used as the bases because of its improved extensibility, effecting feeling and high safety, However, for example, when a passe the orgesses their silicone composition in prepared, it is efficient to obtain a smooth, homogeneous composition in a single system, and low-viscosity silicone oil is easily separated or discharged from the resulting composition, resulting in low setability.

In order to solve the above-mentioned problem of low-viscosity silicone oil, there has been proposed the use of organic materials such as the methods of using fatty acid esters of dectrin as thickners (see Jupanese Patent Unexamined Publication Nos. 52: 121764, 52: 145970, 52: 145971 and 55: 159489), the methods of using fatty acid setter of success (see Jupanese Patent Unexamined Publication No. 52: 253568) and the methods of using interlyality and sophyriviny alcohol or trimetrylelishted polysocharidates (see Jupanese Patent Unexamined Publication No. 62: 46556, 62: 6759 and 63: 17279). However, the use of these organic and inorganic materials as the thickners results in the problem of deteriorating inherent characteristics of low-viscosity silicone is such as referenting feeling and centerbility.

Recently, methods have been proposed in which low-viscosity silicone oil is treated using compounds obtained by partially crossinishing silicones having a specific degree of polymerization as tribeners under shearing torce, thereby obtaining homogeneous paste-like silicone compositions (see Japanese Patent Uhavarnined Publication Nos. 2-42533 and 5-140230). However, these methods require the use of mixers such as ball mills, there on limits and colicid mills having strong shearing force for obtaining paste-like silicone compositions, resulting in disadvartages such as trouble-some preparation, high viscosity if the resulting compositions and restricted compounding amounts of the conditions. Bases have therefore been desired which can make get and increase viscosity by simpler compounding without impairing the internet feeling of the medically evident silicones having a society of 100 or less.

#### SUMMARY OF THE INVENTION

As a result of intensive investigation under such situations, the present inventors have discovered that novel sallcone derivatives in which trizains compounds or printindine compounds are chemically bound to aming open a mino-modified dilicones can gel elicone oil and/or liquid oil or can increase the viscosity thereof simply and stably, thus completing the present invention.

Namely, the present invention provides:

(1) A triazinyl group- or pyrimidinyl group-containing amino-modified silicone derivative having a triazinyl group or a pyrimidinyl group bound to an amino group of an amino-modified silicone, or a derivative in which triazine or pyrimidine is bound to silicone oil.

(2) A triazinyl group- or pyrimidinyl group-containing amino-modified silicone derivative represented by general formula (1), including a silicone derivative in which a triazinyl group or a pyrimidinyl group is chemically bound to silicone od is sepresented by general formula (1) inrespective of its raw material:

$$\begin{array}{c}
N - (CH_1)_m \\
\downarrow \\
R
\end{array}$$

$$\begin{array}{c}
N - (CH_1)_m \\
\downarrow \\
R
\end{array}$$

$$\begin{array}{c}
I \\
Si - O \\
\downarrow \\
R
\end{array}$$

$$\begin{array}{c}
I \\
I \\
R
\end{array}$$

$$\begin{array}{c}
I \\
R$$

$$\begin{array}{c}
I \\
R
\end{array}$$

$$\begin{array}{c}
I \\
R$$

$$\begin{array}{c}
I \\
R
\end{array}$$

$$\begin{array}{c}
I \\
R$$

$$\begin{array}{c}
I \\
R$$

$$\begin{array}{c}
I \\
R
\end{array}$$

$$\begin{array}{c}
I \\
R$$

$$\begin{array}{c}$$

wherein each R, which may be the sum or different, represents hydrogen or a linear, branched-chain or cyclic hydrocathon group of 1 to 8 cathon borns; m and n each represent integer from 1 to 6; he represents an integer from 0 to 6; "a" represents an integer from 1 to 400; and at least two of Y are nitrogen, and the remainder is carbon; (3) The silicone deviative described in the above (1), which is represented by general formula (2), or an armino ellocone derivative in which a teazine ring or a pyrimidine ring is bound to silicone oil having one amino end group at the armine and thereot:

$$\left\{
\begin{array}{c}
N \\
\vdots \\
R
\end{array}
\right\} \left\{
\begin{array}{c}
R \\
\vdots \\
Si \\
R
\end{array}
\right\} \left\{
\begin{array}{c}
R \\
\vdots \\
Si \\
R
\end{array}
\right\}$$
(2)

wherein each R, which may be the same or different, represents hydrogen or a linear, branched-chain or cyclic hydrocarbon group of 1 to 8 carbon atoms; and n each represent integers from 1 to 8.1 represents an integer from 1 to 40; at least two of Y are nitrogen atoms, and the remainder is a carbon atom; prepresents an integer from 1 to 5 in the case of hizariae, or from 1 to 4 in the case of pyrimdine; and when p is 1 or 2 in the case of the presents an integer from 1 to 5 in the case of the presents an integer from 1 to 5 in the case of the presents an integer from 1 to 6 in the case of pyrimdine; and when p is 1 or 2 in the case of pyrimdine; and a high case of pyrimdine; and a pyrimdine and a pyrimdine; and a pyrimdine and pyrimdine and pyrimdine pyrimdine in the case of the pyrimdine in the pyrimdine in pyrimdine in the above (1), which is presented by general formula given and pyrimdine in the present formula by general formula given persent formula the present formula the pyrimdine in the pyr

$$\begin{bmatrix} N & - (CH_{j})_{a} & - (CH_{j})_{a}$$

wherein each R, which may be the same or different, represents hydrogen or a linear, branched-chain or cyclic hydrocarbon group of 11 of 0s actions atoms; in and n each represent integers from 1 to 6; in expresser is an integer from 1 to 40; all least two of Y are intropen atoms, and the remainder is a carbon atom; q is an integer from 1 to 10; p. represents an integer from 1 to 3 in the case of primitines; and when pix is 1 of 2 in the case of brainine, or 1 co 3 in the case of primitines; and when pix is 1 of 2 in the case of brainine, or 1 co 3 in the case of primitines; and when pix is 1 of 2 in the case of brainine, or 1 co 3 in the case of primitines, and when group group, an armino group which may be substituted by a linear or branched alloyd group, of 10 action atoms, or a linear or branched alloyd group, of 10 6 carbon atoms, are all case to branched alloyd group, and the case of branched alloyd group, of 10 action atoms, or a linear or branched alloyd group of 10 action atoms, or a linear or branched alloyd group of 10 action atoms, or a linear or branched alloyd group of 10 action atoms, or a linear or branched alloyd group of 10 action atoms, or a linear or branched alloyd group of 10 action atoms, or a linear or branched alloyd group atoms, are such as a hydrogen atom, may be bound to a residual carbon atom of the triazine ring or the primiting in circ.

# (i) When g is from 1 to 10, A1 and A2 are each represented by general formula (4):

$$-N - (CH_1) = \begin{bmatrix} N - (CH_2) & R & R \\ 1 & Si - 0 & I \\ R & R & R \end{bmatrix} = \begin{bmatrix} CH_2 & N - N \\ 1 & Si - 0 \\ R & R \end{bmatrix} = \begin{bmatrix} CH_2 & N - N \\ R & R \end{bmatrix} = \begin{bmatrix} CH_3 & N - N \\ R & R \end{bmatrix}$$
(4)

wherein R, m, n, h, a and Y have the same meanings as given above; and A1' and A2' each represent hydrogen atoms, hydroxyl groups, amino groups which may be substituted by linear or branched alkyl groups each having 1 to 6 carbon atoms, halogen atoms, carboxyl groups, linear or branched alkyl groups each having 1 to 6 carbon atoms, or linear or branched alkoxyl groups each having 1 to 6 carbon atoms;

or A1 and A2 are each represented by general formula (5):

$$-N - (CH_2)_a = \begin{bmatrix} N - (CH_2)_a & \begin{bmatrix} N \\ 1 \\ 5 - 0 \end{bmatrix} & \begin{bmatrix} N \\ 1 \\ 0 \end{bmatrix} & \begin{bmatrix} N \\ 1 \\ N \end{bmatrix}$$
(5)

# wherein R, m, n, h and a have the same meanings as given above;

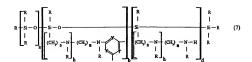
or 11 and A2 each represent hydrogen atoms, hydroxyl groups, amino groups which may be substituted by linear or branched silely groups each having 1 to 6 carbon atoms, halogen atoms, catbody groups, linear or branched allyl groups each having 1 to 6 carbon atoms, or linear or branched alloxyl groups each having 1 to 6 carbon atoms; and

(ii) When q is 0, general formula (3) is represented by general formula (6):

$$\begin{bmatrix} N & \dots & (CH_{r})_{a} & \begin{bmatrix} -N & -(CH_{r})_{a} \\ R & \end{bmatrix} \begin{pmatrix} R & 1 \\ Si & -0 \\ R & a \end{bmatrix} & \begin{bmatrix} R & -(CH_{r})_{a} & -N \\ R & R \end{bmatrix} \begin{pmatrix} CH_{r} & -N \\ R & R \end{bmatrix} \begin{pmatrix} CH_{r} & -N \\ R & R \end{pmatrix} \begin{pmatrix} CH_{r} & -N \\ R & R \end{pmatrix} \begin{pmatrix} CH_{r} & -N \\ R & R \end{pmatrix}$$

said silicone derivative inclusing a derivative in which 1 to 3 carbon atoms of the triazine ring or 1 to 4 carbon atoms of the primitine ring are bound to a\_v-diamine instance of at both arising near the restriction of the structure composed of 2 to 10 repeating units thereof which includes a linear, branched and/or network structure, and including a derivative in which one end of the a\_v-diamine silicone of lamp be an animing group or an animo group substituted by a linear or branched alleyl group of 1 to 6 carbon atoms without bonding to the tri-axing group or the primitivity group.

(5) The silicone derivative described in the above (1) represented by general formula (7), which is obtained by allowing a side-chain amino silicone oil to react with a reactive group-containing triazine or pyrimidine compound:



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wherein each R, which may be the same or different, represents hydrogen or a linear, branched-chain or cyclic hydrocarbon group of 1 to 8 carbon atoms; m and n each represent integers from 1 to 6; h represents an integer from 0 to 6; "a" and "b" each represent integers of 1 or more, "d" represents an integer of 0 or more, and "a-b-d" represents an integer of 0 or less; "a", "b" and "d" each show the ratios, and do not specify the order of arrangement, and carbon to or "ye remainder is a carbon atom;"

(6) A method for producing the silicone derivative described in the above (1) which comprises allowing a amino silicone oil to react with a reactive group-containing triazine or pyrimidine compound;

(7) The method described in the above (6), in which the amino silicone oil is represented by the following general formula (8):

$$B - \begin{cases} I & R \\ Si - O \end{bmatrix} = \begin{cases} R & R \\ Si - O \end{bmatrix} = \begin{cases} R & Si - B^* \end{cases}$$
 (8)

wherein each R, which may be the same or different, represents hydrogen or a linear, branched-chain or cyclic hydrocathon group of 1 to 8 carbon atoms; a and b each represent integers of 1 or more, and the sum of a and b represents an integer of 400 or less; a and b each show the ratice, and do not specify the order of arrangement; and at least one of 8, B° and B° is a group represented by general formula (9):

wherein each R', which may be the same or different, represents hydrogen or a linear, branched or cyclic hydrocarbon group of 1 to 8 carbon atoms; m and n each represent integers from 1 to 6; and h represents an integer from 0 to 6: and

the reactive group-containing triazine or pyrimidine compound is a compound represented by the following general formula (10):

15 wherein at least one of X<sup>1</sup>, X<sup>2</sup> and X<sup>3</sup> represents a halogen atom, the remainders each represent hydrogen atoms, hydroxyl groups, amino groups which may be substituted by linear or branched allkyl groups each harwing 1 to 6 carbon atoms, carboxyl groups, linear or branched allkyl groups each harwing 1 to 6 carbon atoms; or linear or branched allkyl groups each harwing 1 to 6 carbon atoms; and at least two of Y are nitrogen atoms, and the other is a carbon atoms.

 (8) A gelling agent comprising the triazinyl group- or pyrimidinyl group-containing silicone derivative described in the above (1), (2), (3), (4) or (5);

(9) A composition comprising the triazinyl group- or pyrimidinyl group-containing silicone derivative described in the above (1), (2), (3), (4) or (5) and silicone oil and/or liquid oil; and (10) A base comprising the triazinyl group- or pyrimidinyl group-containing silicone derivative described in the

above (1), (2), (3), (4) or (5).

# DESCRIPTION OF THE PREFERRED EMBODIMENTS

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# The present invention will hereinafter be described in detail.

When the novel allicone derivatives of the present invention are produced, in addition to methods in which arrindosticone oils allowed to react with reactive group-containing triazine or primitine compounds are new materials are methods can be employed in which reactive group-modified eliicones having reactive groups such as halogen and arrind proup-containing traizine or pryimidine compounds are allowed to react with each other, which has a reversed relationship in the reactive groups with the first method. This is the reason why the sitions derivatives of the present invention include all compounds represented by general brank (1). However, the reactivity of the nearly groups such as habgen on heterocyclic molety is stronge in the first method, so that the first method is generally used. The first method is hereinafter manife described but the second method can also be used strainlar.

As the starting materials for producing the novel silicone derivatives of the present invention, the case of using amino silicones is lirist described. The sites of amino modification may be any of side chains, one end and both ends, 40 and the number of the amino groups, the position thereof, etc. are not limited.

These amino silicones are represented by the following general formula (11):

$$\begin{array}{c|c}
R & R \\
B - Si - O & Si - O \\
I & R & R
\end{array}$$

$$\begin{array}{c|c}
R & R \\
I & Si - O \\
I & I \\
B' & R
\end{array}$$

$$\begin{array}{c|c}
R & R \\
I & Si - B'' \\
I & R
\end{array}$$
(11)

wherein each R, which may be the same or different, represents a linear, branched-chain or cyclic hydrocarbon group
of 1 to 8 carbon atoms; at least one of B, B' and B" is a group represented by general formula (12):

wherein each Pf, which may be the same or different, represents hydrogen or a linear, transhed or cycle hydrocarbon group of 1 to 8 carbon atoms, m and n each represent integers from 1 to 6, and h represents an integer from 0 to 6; the remainders of 8, 8° and 8° each represent linear, branched-chain or cyclic hydrocarbon groups each having 1 to 8 carbon atoms; a and b each represent integers of 1 or more, and the sum of a and b ender persents an integer of 400 or less; and a and be ach show the ratios, and do not specify the order of amanquement.

Examples thereof include amino silicones having one amino end group represented by the following general formula (13):

we wherein each R, which may be the same or different, represents hydrogen or a linear, branched-chain or cyclic hydro-carbon group of 1 at 5 carbon atoms, B represents a group represented by general formula (12); and a represents an integer from 1 to 400;

a,a-diamino silicones represented by the following general formula (14):

s wherein each R, which may be the same or different, represents hydrogen or a linear, branched-chain or cyclic hydro-carbon group of 1 to 8 carbon atoms; B and B' each represent groups represented by general formula (12); and a represent sa mitteger from 1 to 400; and

side-chain amino silicones represented by the following general formula (15):

wherein each R, which may be the same of different, represents hydrogen or a linear, branched-chain or cyclic hydrocation orgood of 10 darbot atterns. B represents a goog represented by general formula (12); and the each depresent integers of 1 one one, and the sum of and b represents an integer of 400 or less; and a and b each show the redisc, and do not society the order of armenement.

In the compounds represented by general formulas (13), (14) and (15), the amino silicone derivatives used in the present invention, examples of the hydrocarbor groups seach having 1 to a carbon atoms represented by it includes inear hydrocarbon groups such as methy, ethyl, proxyl, bally, pentyl, heryl, heryl, and ochly, branched hydrocarbon groups such as isopropial and isolancy, allendy groups cauch as virty and tally, cycleding groups such as cyclotheoly and cyclotheoly and any groups such as specification, and tally the composition of the cyclotheoly and cyclotheoly and any groups such as specification as a facility of the cyclotheoly and cycl

10 and examples thereof include 3,3,3-trifluoropropyl. Methyl and phenyl are preferred among others, and methyl is particularly preferred.

In the compounds represented by general formulas (13), (14) and (15), the number of silicon atoms contained in a modified silicone molecule (a+b) is 2 to 400, preferably 10 to 300, and more preferably 40 to 150. Examples of the armino silicones used in the present invention include the following:

Amino Silicone Oil Manufactured by Dow Corning Toray Silicone Co., Ltd.

Structure (1) (a, w-diamino silicone)

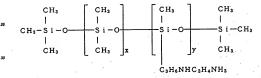
Structure (2) (Side-Chain amino silicone)

R: Alkylene

ED 0 764 470 82

Product Name	Viscosity (cs)	Specific Gravity	Refractive Index	Remarks	NH <sub>2</sub> equivalent
BY16-853	30	0.96	1.414	α,ω-diamino silicone	650
BY16-853B	80	0.97	1.407	ditto	2200
BY16-828	120	1.05	1.453	side-chain amino silicone	3500
BY16-850	1100	0.98	1.411	ditto	4000
SF8417	1200	1.07	1.452	ditto	1800
BY16-849	1300	1.03	1.426	ditto	.600
BY16-872	18000	1.01	1.421	ditto	2000
BX16-755B	240	١.	1.408	secondary amine	2200
BX16-193				one amino end group	4000

# Amino Silicone Oil Manufactured by Nippon Unicar Co., Ltd.



Product Name	Amino Equivalent	Si Polymerization Degree
FZ-3705	4,000	140
FZ-3707	1,400	50
FZ-3710	1,700	300
FZ-3712	1,700	200

mino Silicone Oil Manufactured by Dow Corning Toray Silicone Co., Ltd.

Product Name	Appearance	Viscosity (cs)	Specific Gravity	Refract- ive Index	Amino Eq- uivalent		
SF8417	light yellowish brown, transparent	1200	96.0	1.407	1800	CH, CH, ÇH	ÇH, ÇH,
BY16-828	light yellow, transparent	120	96.0	1.408	3500	-	0), \$ 1 CF
BY16-849	ditto	1300	0.99	1.414	009	X Z	_
BY16-850	ditto	1100	0.97	1.405	4000	. r ×−z	
BY16-872	ditto	18000	0.98	1.407	2000		
BY16-853	light yellow, transparent	30	96.0	1.414	650	ен, сн.	
BY16-853B	colorless, trans- parent	08	0.97	1.407	2200	H,NR(\$10).SIRNH, CH, CH,	r Z
BX16-755B		240		1.408			ı

The molecular weight of these amino silicones is 100 to 30,000, preferably 1,000 to 20,000, and more preferably 3,000 to 10,000.

The triazine and pyrimidine compounds having reactive functional groups which are used for reaction with the amino silicones are represented by the following general formula (10):

ss wherein at least one of X1, X<sup>2</sup> and X<sup>3</sup> represents a hatogen atom, the remainders each represent hydrogen atoms, hydroxyl groups, aminor groups which may be substituted by linear or branched schain alloy groups each having 1 to 6 carbon atoms, carboxyl groups, linear or branched alloxyl groups each having 1 to 6 action atoms, and a least word V1 are nifrogen, and he other is carbon.

The reactive groups of these triazine or symindrine compounds may be any, as long as they are groups which reads with aming oppose of the amino silicones. Examples hereof include histogran atoms such as fluorine, obtains, bronne and lodine. Although substituted groups XI and XI of the triazine skeleton may also be any, examples thereof include hydrogen, hydrooy, amino groups which may be substituted by ally groups, habgen, carbony, linear or branched alloy groups each having 1 to 6 carbon atoms, or linear or branched alloy groups each having 1 to 6 carbon atoms, or linear or branched alloy groups each having 1 to 6 carbon atoms, or linear or branched alloy groups each having 1 to 6 carbon atoms examples of the alloyi-dusbifield and groups include elipharinin, propylamino and isopropylamino. Further, the carbonyl groups contain salts thereof such as allottine salts and alkaline earth metal salts, and examples thereof include softum, potassium, calinium, magnesium, aluminum and zinc salts. Examples of the linear or branched alloy groups such as metally, ethyl, propyl, butyl, pertyl and heptyl, branched alloy groups. Such as sopropyl, isochly, soperally and isopropy, and those of logical alloy groups such as open atoms include filment box, ethony, proposity, butylon, pertyl and polytogroups and bytogroups and hereofron, ethony, proposity, becompay and othershow, ethors, proposy, button, pertalsen, becopposy and dimethou.

Examples of these iriazine compounds include operative cholde, 2-chloro-4,6-bis(methylamino)-5-triazine, 2-chloro-4,6-bis(mylamino)-5-triazine (simazine), 2-chloro-4,6-bis(proylamino)-5-triazine, 2-chloro-4,6-bis(proylamino)-5-triazine, 2-chloro-4-methylamino-6-triazine, 2-chloro-4-methylamino-6-triazine, 2-chloro-4-triaylamino-6-triazine, 2-chloro-4-triylamino-6-triazine, 2-chloro-4-triylamino-6-triylamino-6-triylamino-6-triazine, 2-chloro-4-triylamino-6-triylamino-6-triazine, 2-chloro-4-triylamino-6-triylamino-6-triylamino-6-triylamino-6-triylamino-6-triylamino-6-triylamino-6-triylamino-6-triylamino-6-triylam

Examples of the pyrimidine compounds include 2-chloropyrimidine, 2-chloro-4,6-dimethylpyrimidine, 2-chloro-4,6-dimethylpyrimidine, 2-chloro-4,6-dimethylpyrimidine, 2,5-dichloro-4-methoxy-6-trichloromethylpyrimidine and 2,4,5,6-tetrachloropyrimidine.

On the other hand, in addition to the methods in which the above-mentional armino silicones are allowed to react with the reactive group-containing trizative or primition ecompounds as starting materials, the novel silicone deviatives of the present invention can also be produced by allowing reachine group-containing trization or primition ecompounds. When the reactive group-containing silicones for react with armino group-containing trization or primition for ecompounds. When the reactive group-containing silicones or used, halpons printing to the reactive groups. The halpons silicones induce, for example, other silicones oil and thosy silicones oil include 2.4.6 triamino-6-trizative (melarine), 2.4-diamino-6-dimethyl-fristative and 2.4-diamino-6-dimethyl-fristative and 2.4-diamino-6-dimethyl-gradienty

Reaction solverts used in the reaction of these amino silicones with the triazine or pyrimidine compounds or in the reaction of the halopen silicone oil with the amino group-containing triazine or pyrimidine compounds may be any, as long as they dissolve both of them. Such solvents include between, tokere, sylien, n-haxane, n-heptane, carbon tel-rachfords, chrotrom, perchloredtylens, tiptichoredtylens and chloroberaren.

The reaction is conducted by dissolving the amino silicone oil or the halogen silicone oil and the triazine compounds or the pyrimidine compounds in the reaction solvents, and heating and adding catalysts if necessary.

The reaction is conducted at norm temperature or higher, preferably at 70 to 120°C, and basic compounds such as pyridine, artificial and alkylamines can be used as the reaction catalysts. After the reaction, unreacted halogen groups are substituted by water, alcohols, etc., followed by solvent removal and removal or furneacted products. Then, the resulting products are subjected to vacuum drying to obtain the novel silicone derivatives in which triazinyl groups or pyrimidinyl groups are bound to the arrino silicone oil or the halogen silicone oil.

The properties such as gel strength and viscosity can be controlled by changing the binding ratio of the amino silicone to the triazinyl group or the pyrimidinyl group.

5 For example, when added to liquid oil or silicone oil generally used for cosmetic or pharmaceutical preparations, a disubstituted compound in which silicone oil is introduced into two positions of the firstiant young or the primidityry group terms a hard pel, a mono-substituted compound in which silicone oil is introduced into oil three positions gives a weak get or a viscoshy-increased fluid liquid. With respect to cometics, for example, the disabstituted compounds are used to foundate tooks, employed and the production of the compounds are used to foundate to compounds. The production of the production. Further, the disabstituted compounds are used for celeration and electronic parts and vibration absorbing agents for almorphism, and the threubstituted compounds are used for thickness, sealing agents and pigment dispersing agents. Thus, any one of the mono-substituted, disubstituted and tri-substituted compounds are used for thickness.

Reaction products in which halogen is left cause initiating feeling when they achieve to the skin. They are therefore unsuitable for cosmetics and pharmaceutical preparations. It is therefore preferred that halogen is substituted as described above. However, the reaction products containing halogen can be used for other applications such as water repelling agents, potting agents for electric and electronic parts and vibration absorbing agents for dampers, etc. without any particular problem.

When the armine silicone is an  $\alpha_i$  ordinative silicone or a side chain armino silicone, a compound in which triazinyl or pyrimiding lygous are bound to all armino gougo of the  $\alpha_i$ -distrince silicone or the side chain armino silicone of the discholar in silicone of the side of the animal silicone of the side of the silicone of the si

For example, when the side chain amino silicone has amino groups at 40 positions in a molecule, a gelling agent can be obtained by introducing history for pyrimidinyl groups into 2 or 3 positions, said gelling agent giving a homogeneous, flexible, soft gel when added to silicone oil having a viscosity of 100 ce or less.

The compounds of the present invention represented by general formula (3) which are obtained by the reaction of the a\_ordinant solicones with reactive fruitaine or primitine can be compounds showing full(b), compounds showing individually increasing properties, gel-like compounds or nubber-like elestomers, depending on the reaction conditions such as the reaction tomizer and the reaction time.

For example, when a soft get is desired, the reaction time is shortened, or the amount of the reactive group-comtaining trizine or pyrimidine compound in reduced in the reaction. When a harder get is desired, the reaction time is protonged, or the amount of the reactive group-containing triazine or pyrimidine compound is increased. Thus, the hardness of roduct cells can be controlled.

These compounds vary in viscosity according to the degree of polymerization (molecular weight) represented by q in general formal (8), and a higher molecular weight results in a higher viscosity, in addition to this, the linear structure and the crosslinked structure are obtained as the molecular structure according to methods of polymerization. The first estructure results in soleter compounds compared with the crosslinked structure. For example, get-filia compounds rich in the linear structure restructure rear soft, whereas gets rich in the crosslinked structure become hard, resulting in elastic, rub-er-filia commounds.

Specifically, the products are described as follows.

When the products are mainly composed of the compounds represented by general formula (3) wherein q is 3 or less, the products are poor in the crossifixed structure and rich in the ineval structure. Dissolution thereof in silicotion results in various compounds extending from high viscous materials exhibiting viscosity increasing properties to week and like commounds.

When the products are mainly composed of the compounds represented by general formula (3) wherein q is 5 to 50, the crosslinked structure is nich, resulting in exhibition of properties extending from weak gel-like properties to hard nither-like properties.

When the products are mainly composed of the compounds represented by general formula (3) wherein q exceeds 10, the products are almost all composed of the crosslinked structure and further composed of the polymer structure, resulting in exhibition of properties extending from rubber-like properties to resh-filke properties.

Similarly to the quediamins silicones, the compounds of the present invention represented by general formula (7) which are obtained by the reaction of the side-chain amino silicones with reactive triazine or pyrimidine compounds vary in viscosity according to the degree of polymerization (molecular weight) represented by (a-b-d), and a higher molecular weight results in a higher viscosity. For the linear structure and the crosslinked structure developed as the molecular structure accordant to methods of cohvenization. The linear structure is results in softer compounds command with

the crosslinked structure. For example, gel-like compounds rich in the linear structure are flexible and soft, whereas gels nich in the crosslinked structure become hard, resulting in elastic, nubber-like compounds.

When cyanuric chioride is used as the triazine compound, it can bind to reache groups of the allicone at a maximum of 3 positions, because it has 3 reachey copys. The structure of the sictions obtained heady is ago to the test or considered structure, compared with a compound having one reache group such as 2-chloropyrimidine. The resulting compound has therefore to extend the observable such

Further, more amino groups in the molecule of the amino silicone results in more reactive positions, so that the resulting compound is apt to take the crosslinked structure, leading to a hard compound. Accordingly, the use of the silicone having a small amount of amino groups can provide the silicone derivative having desired hardness and proper-

an example of the ppoffic structure, the side-chain amino silicone is got to take the intermolecular or intramolecular crosslinked structure when the triazine or pyrimidine compound having at least 2 reactive groups is used, resulting in a hard compound. In order to avoid this problem, the introduction rate of the triazine or pyrimidine compound is decreased, or the triazine compound having one reactive group such as 2-chirors.—Belieflyeliminoly—fritizarie or the pyrimidine compound such as 2-chiroryprimidine is used, wheetey a compound having moderate get properties can be

More amino groups in the side-chain amino silicone oil results in a harder get. It is therefore preferred that the number of amino groups is proper. In order to obtain the soft get-like product, the number of amino groups in the molecule is preferably for less.

As described above, the hard gels, the soft gels and the liquid triazinyl group- or pyrimidinyl group-containing amino silone derivatives are obtained according to various conditions, and they can be selected depending on each application, similarly to the above-mentioned various substituted compounds.

The novel triazinyl group- or pyrimidinyl group-containing amino silicone derivatives of the present invention are dispersed in liquid oil or silicone oil, and dissolved by heating, billowed by cooling it necessary, thereby obtaining compositions extending from oil-file compositions to viscous materials:

The amount of the taining froup- or perimiding group-containing amino allicone deviative of the present invention added to the allicinone oil varies with the molecular weight, etc. of the triasing group- or primitingly group-containing amino silicone derivative, and a higher molecular weight and a higher viscosity of the allicone derivative require a smaller amount thereof added. On the other hand, a lower molecular veryint and a lower viscosity of the allicone derivative require a larger amount thereof added. In the latter case, the triazing group- or primitingly group-containing amino silicone derivative titled can also be used as a base. When used as a gailing agent, the triazing group- or primitingly group-containing amino silicone derivative titled can also be used as a base. When used as a gailing agent if the present invention is added generally in an amount of 3% by weight or more, any primiting group- or group- or

The liquid oil may be any as long as it is compatible with the silicone derivatives of the present invention, and examples thereof include hydrocarbon oil such as isoparatifin oil, and ester oil. The silicone oil may be any as long as it is liquid, and chain sicones, cyclic silicones, arinor, cathonyl- or lookhot-silicones, etc. can be used.

The triazinyl group- or pyrimidinyl group-containing amino silicone derivatives of the present invention are dissolved in low-viscosity silicone oil, hydrocatoon oil, polar oil, et., to cause gelation or an increase in viscosity. They can be therefore used as bease for various industrial materials as well as for committee and pharmaceutical preparations. In addition, the triazinyl group- or pyrimidinyl group-containing amino silicone derivatives of the present invention themselves can be used as bases for commetcs, pharmaceutical preparations and various industrial materials.

When, for example, silicone oil having a low viscosity of 100 cs or less generally used in cosmetics and pharmaceutical preparations is used as a soft, homogeneous get, the use of the mono-abstituted compound in which a triactny group or a pyrindidity group of the amino silicone is substituted provides a get-like composition not impairing refreshing feeling of the silicone oil. When used as a hard, homogeneous get high in form retention, the di-substituted compound is mainly emolypoid.

As described above, when the silicone derivatives are used in cosmetics and pharmaceutical preparations, it is preso ferred that halogen is removed by substitution, etc.

The silicone oil is used for various industrial materials utilizing its insulating properties and water repellency, and can be used properly by various selections according to its purpose of use as a gelling agent.

The gel-like compositions comprising the triazinyl group- or pyrimidinyl group-containing amino silicone derivatives of the present invention and silicone oil are thermally reversible, and addition of DMSO to the gel-like compositions results in fluid liquids. From this fact, this gelation is deduced to be mainty caused by a hydrogen bond.

The present invention will be described in more detail with the following examples. It is understood of course that they are for the purpose of illustration only and are not intended to limit the scope of the invention.

# EXAMPLE 1

Three grams (1.2X10<sup>2</sup> mol) of a side-chain amino silcone (BY16-849 manufactured by Dow Coming Toray Silcone Co., Ltd., MW: ca. 25000, amino equivalent: 600) dissolved in 20 g of benzene was mixed with 0.57 g (s.0X10<sup>2</sup> mol) of 2-dhotopyrindrine previously dissolved in 20 g of benzene, and the mixture was stirred under reflux for 24 hours. After termination of the reaction, the solvent was removed by distillation under reduced pressure to obtain a 2prindrindrivoranopolysilozane within his priminding vigrous bound.

# EXAMPLE 2

Thirty grams (6.6X10<sup>3</sup> mol) of a both-terminal amino ellicone (BY16-83SI manufactured by Dow Coming Tony Silicone Co., Ltd., MW: ca.4500, amino equinalent: 2250) dissolved in 15 g of benzene was mixed with 2.46 g (13.3X10<sup>3</sup> mol) of cyanutic chlorida previously dissolved in 15 g of benzene, and the mixture was stirred at room temperature for 24 hours. After termination of reaction, the solvent was removed by distillation under reduced pressure, blowed by seashing with water. The resulting product was heated again under reduced pressure to memore the solvent by distillation, thereby mainly obtaining a triaziny/organopolysiloxane in which triazinyl groups were bound to both ends. This product showed the following nuclear magnetic resonance specia:

Table 1

Structure Confirmation According to Nuclear Magnetic Resonance Spectra (13C-NMR)

Structure	Chemical Shift 8
$\frac{1}{C}$ -NH-R-Si(R) <sub>3</sub>	166.1 ppm
[- c-c1	169.9 ppm 171.1 ppm
Solvent: CDCl,	

#### EXAMPLE 3

3.1 g. (1.24XiO<sup>4</sup> mo) of a side-chain anino elifono (BY16-84) manufactured by Dow Coming Tomy Silicone Ca, Ltd, MW: ca. 25000, amino equivalent: 600) dissolved in 50 g of 1.4-dioxane was mixed with 0.1 g (4.95XiO<sup>4</sup> mol) of 2-chioro-4,6-bis[ethylamino]-e-triazine previously dissolved in 50 g of 1.4-dioxane, and the mixture was selimed under retux at the boiling point for 24 hours. After termination of the reaction, the solvent was removed by desiliation under reduxed pressure to mainly obtain a 24,46-bis[ethylamino]-e-triazing/oraprojs-biscane in which 2-chioro-4,6-bis[ethylamino]-e-triazing groups were bound to a part of amino groups of one molecule of the side-chain amino silicone. This product showed the following nuclear imagnific resonance specta:

Table 2

Structure Confirmation According to Nuclear Magnetic Resonance Spectra ( $^{13}\mathrm{C-NMR}$ )

Structure	Chemical Shift δ
$-\underline{C}H_3$ (of an aminoethyl group)	15 ppm
$-\underline{C}H_2$ - (of an aminoethyl group)	35 ppm
$\frac{1}{C}$ -NH-R-Si(R) <sub>3</sub>	163 ppm
(carbon of a triazinyl ring to which a silicone was bound)	. *
C-NH-CH <sub>2</sub> -CH <sub>3</sub>	165 ppm
(carbon of a triazinyl ring to which an amino- ethyl group was bound)	
Solvent: CDCl <sub>3</sub>	

# EXAMPLE 4

35 11.19 g (2.48X10<sup>-3</sup> mol) of an c.,a-diamino silicone (BY16-4848 manufactured by Dow Coming Toray Silicone Co. Ltd., MW. c. at 650, mirno equivalent :2500) dissolved in 150 g of 1.4-dioxane, and the mixture was refluxed for 2d-chloro-4,8-bis(ethylarmino)-4-triazine previously dissolved in 150 g of 1.4-dioxane, and the mixture was refluxed for 2d-hours with siming at 50°C. Alter termination of the reaction, the solvent was removed by distillation under reduced pressure to obtain a 2-14,6-bis(ethylarmino)-1-triaziny/ groups were bound to both ends. This compound was highly visious and repate-like.

# EXAMPLE 5

0.22 g (1.19X10<sup>-9</sup> mol) of cyanuric chloride dissolved in 20 g of berezene was mixed with 5.0 g (1.11X10<sup>-9</sup> mol) of silicone oil having one amino end group (MM: ca. 4500), followed by stirring for 24 hours. After termination of the reaction, the solvent was removed by distillation under reduced pressure to mainly obtain a triazinylorganopolysiloxane in which one silicone chain was bound to a triazinyl group.

#### EXAMPLE 6

0.11 g (0.58X10<sup>3</sup> mol) of cyanuric chloride dissolved in 20 g of berzene was mixed with 5.0 g (1.11X10<sup>3</sup> mol) of silicone oil having one armino end group, 8X16-193, billowed by silimig for 24 hours. After termination of the residence, the solvent was removed by distillation under reduced pressure to mainly obtain a triazinytorganopolyaloxane in which two silicone chains were bound to a triazine compound.

### EXAMPLE 7

0.07 g (0.37X10<sup>-3</sup> mol) of cyanuric chloride dissolved in 20 g of benzene was mixed with 5.0 g (1.11X10<sup>-3</sup> mol) of silicone oil having one amino end group, BX16-193, followed by stirring for 24 hours. After termination of the reaction,

the solvent was removed by distillation under reduced pressure to mainly obtain a triazinylorganopolysitoxane in which three silicone chains were bound to a triazine compound.

#### EXAMPLE 8

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Amino silicone derivatives (1) to (7) obtained in Examples 1 to 7 were each added to respective silicones so as to give a concentration of 10% by weight or 20% by weight, and dissolved by heating at room temperature or 80°C with stirring. The resulting solutions were each allowed to stand at room temperature, followed by observation. The gelling and viscosity increasing properties of the derivatives obtained in Examples 1 to 7 to the silicones are shown below.

Sample Concentration (%)		Silicone		
		Silicone		
	No. 1	No. 2	No. 3	
Example 1 10	Δ	Δ	Δ	
. 20	0	0	0	
Example 2 10	0	0	0	
20	0	0	- 🚳	
Example 3 10	Δ	Δ.	Δ .	
20	Δ.	Δ		
Example 4 10	Δ	Δ	Δ	
20	Δ.	Δ.	Δ	
Example 5 10	0	0	Δ	
20	0	0	Δ	
Example 6 10	0	0	0	
20	0	0	0	
Example 7 10	0	0	Δ	
20	0	0	Δ	
No. 1: dimethylpolysiloxane 5 cs No. 2: dimethylpolysiloxane 50 cs				

@: gel (form retention kept)

O: weak gel

A: increase in viscosity

X: solution

The equivalent amount of DMSO was added to each get of No. 1 obtained in Example 1, 2, 5, 6 or 7 shown in Table 3, followed by stirring. As a result, each get was turned solution-like. From this fact, this get was deduced to be mainly made by a hydrogen bond.

The novel silicone derivatives of the present invention can be easily synthesized from the amino silicones and the triazine or pyrimidine compounds, and can gel silicone oil or increase the viscosity thereof stably and homogeneously. The gelled products are thermally reversible, and easily produced. Further, the resulting gels are transparent. The reason for this is deduced to be that this gel is mainly made by a hydrogen bond.

#### Claims

- 1. A triazinyl group- or pyrimidinyl group-containing amino silicone derivative having a triazinyl group or a pyrimidinyl group bound to an amino group of an amino silicone.
  - 2. A triazinyl group- or pyrimidinyl group-containing amino silicone derivative represented by general formula (1):

$$N - (CH_j)_{n} = \begin{bmatrix} N - (CH_j)_{n} & \vdots \\ N - (CH_j)_{n} & \vdots \\ N - (CH_j)_{n} & \vdots \end{bmatrix}$$
(1)

wherein each R, which may be the same or different, represents hydrogen or a linear, branched-chain or cyclic hydrocarbon group of 1 to 8 carbon atoms; m and n each represent integers from 1 to 8; it presents an integer from 0 to 5; "represents an integer from 0 to 5;" represents an integer from 1 to 400; and at least two of Y are nitrogen, and the remainder is carbon.

3. The silicone derivative according to claim 1, which is represented by general formula (2):

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$$\left\{
\begin{array}{c}
N - (CH_{i})_{n} \\
\vdots \\
R
\end{array}
\right\}
\left\{
\begin{array}{c}
R \\
\vdots \\
Si - O
\end{array}
\right\}
\left\{
\begin{array}{c}
R \\
\vdots \\
Si - R
\end{array}
\right\}$$
(2)

wherein each R, which may be the same or different, represents hydrogen or a linear, branched-chain or cyclic hydrocarbon group of 1 to 8 carbon atoms; m and n each represent integers from 1 to 6; h represents an integer from 0 to 6; "at represents an integer from 1 to 40; at least two of Y are integer atoms, and the reniander is a carbon atom; p represents an integer from 1 to 3 in the case of briazine, or from 1 to 4 in the case of pyrimidine; and when p is 1 of 2 in the case of triazine, or 1, 2 or 3 in the case of pyrimidine, a hydroxy group, an entino group which may be substituted by a linear or transched alsky group of 1 to 6 carbon atoms, a halogen atom, a carbonyl group, a linear or branched alsky group of 1 to 6 carbon atoms, or a linear or branched alloxy group of 1 to 6 carbon atoms, as swell as a hydrogen atom, may be bound to a residual carbon atom of the triazine ring or the pyrimidine ring.

40 4. The silicone derivative according to claim 1, which is represented by general formula (3):

$$\begin{bmatrix} N - (CH_{j})_{a} & \begin{bmatrix} 1 \\ R \end{bmatrix} & CH_{j} \\ R \end{bmatrix} \begin{bmatrix} R \\ 1 \\ SI - O \end{bmatrix}_{a} \begin{bmatrix} R \\ 1 \\ R \end{bmatrix} \begin{bmatrix} CH_{j} \\$$

wherein each R, which may be the same or different, represents hydrogen or a linear, branched-chain or cydic hydrocarbon group of 1 to 8 cathor latms; m and neach represent insignes from 1 to 5 in presents an integer from 1 to 400; at least two of Y are nitrogen, and the remainder is carbon; q represents an integer from 1 to 400; at least two of Y are nitrogen, and the remainder is carbon; q represents an integer from 5 to 1 to; prepresents an integer from 1 to 3 in the case of primidine, and when p is 1 or 2 in the case of trainine, or 1,2 or 3 in the case of primidine, and when pot a for 2 in the case of primidine, and when pot and the case of primidine, and when pot a form of the case of primidine, and when pot a form of the case of primidine, and when pot the case of primidine, and when pot the case of primidine, and when potentially group of 1 to 6 cannot along, or a linear or branched alloys.

group of 1 to 6 carbon atoms, as well as a hydrogen atom, may be bound to a residual carbon atom of the triazine ring or the pyrimidine ring;

(i) when g is from 1 to 10, A1 and A2 are each represented by general formula (4):

$$-N - (CH_{2}) = \begin{bmatrix} 1 & CH_{2} & 1 & 1 \\ -N & (CH_{2}) & 1 & 1 \\ -N & N \end{bmatrix} \begin{bmatrix} R & 1 & 1 \\ -N & N \end{bmatrix} \begin{bmatrix} R & 1 & 1 \\ -N & N \end{bmatrix} \begin{bmatrix} CH_{2} & -N & 1 \\ -N & N \end{bmatrix} \begin{bmatrix} CH_{2} & -N & 1 \\ -N & N \end{bmatrix} \begin{bmatrix} CH_{2} & -N & 1 \\ -N & N \end{bmatrix} \begin{bmatrix} A^{1} & 1 & 1 \\ -N & N \end{bmatrix} \begin{bmatrix} A^{2} & 1 & 1 \\ -N & N \end{bmatrix}$$
(4)

wherein R, m, n, h, a and Y have the same meanings as given above; and A1' and A2' each represent hydrogen atoms, hydroxyl groups, amino groups which may be substituted by linear or branched alkyl groups each having 1 to 6 carbon atoms, halogen atoms, carboxyl groups, linear or branched alkyl groups each having 1 to 6 carbon atoms, or linear or branched alkoxyl groups each having 1 to 6 carbon atoms; or A1 and A2 are each represented by general formula (5):

wherein R, m, n, h and "a" have the same meanings as given above;

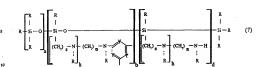
or A1 and A2 each represent hydrogen atoms, hydroxyl groups, amino groups which may be substituted by linear or branched alkyl groups each having 1 to 6 carbon atoms, halogen atoms, carboxyl groups, linear or branched alkyl groups each having 1 to 6 carbon atoms, or linear or branched alkoxyl groups each having 1 to 6 carbon atoms; and

(ii) when q is 0, general formula (3) is represented by general formula (6):

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$$\begin{bmatrix} N & C(H_1) \\ N & C(H_2) \end{bmatrix}_a \begin{bmatrix} N & C(H_2) \\ N & N \end{bmatrix}_a \begin{bmatrix} N \\ N \\ N \end{bmatrix}_a \begin{bmatrix} N \\$$

The silicone derivative according to claim 1 represented by general formula (7), which is obtained by allowing a side-chain amino silicone to react with a reactive group-containing triazine or pyrimidine compound:



wherein each R, which may be the same or different, represents hydrogen or a linear, branched chain or cyclic hydrocarbon group of 1 to 8 carbon atoms; in and n each represent integers from 1 to 6, in represent an integer from 0 to 6; "a "and "b' each represent integers of 1 or none, "of represents an integer of 400 or less; "a", "b" and "of each show the ratios, and do not specify the order of arrangement; and el less the of 1 ye in ribonen atoms, and the remainder is a carbon atom.

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- A method for producing the silicone derivative according to claim 1 which comprises allowing an amino silicone oil to react with a reactive group-containing triazine or pyrimidine compound.
- 7. The method according to claim 1, in which the amino sificone oil is represented by the following general formula (8):

wherein such R, which may be the same or different, represents hydrogen or a linear, branched-chain or cycle hydrocachon group of 1 to 8 cathon alters, "a" and "b" cheach represent integers of 1 or more, and the sum of "a" and "b" represents an integer of 400 or less; "a" and "b" each show the ratios, and do not specify the order of arrangement; and all less to not 9 B, B and "b" is a group, presented by general formula (9):

$$\begin{bmatrix}
(CH_1) & -N \\
 & | \\
 & | \\
 & R'
\end{bmatrix}$$
(CH<sub>1</sub>)  $-N - H$ 
(9)

wherein each R', which may be the same or different, represents hydrogen or a linear, branched or cyclic hydrocarbon group of 1 to 8 carbon atoms; m and n each represent integers from 1 to 6; and h represents an integer from 0 to 6; and

the reactive group-containing triazine or pyrimidine compound is a compound represented by the following general formula (10):

- wherein at least one of X<sup>1</sup>, X<sup>2</sup> and X<sup>3</sup> represents a halogen atom, the remainders each represent hydrogen atoms, hydrody groups, amino groups which may be substituted by linear or branched alloy groups each having 1 to 6 carbon atoms, activory groups, linear or branched alloy groups each having 1 to 6 carbon atoms, or linear or branched alloyd groups each having 1 to 6 carbon atoms; and at least two of Y are nitrogen atoms, and the other is a carbon of the carbon.
- A gelling agent comprising the triazinyl group- or pyrimidinyl group-containing silicone derivative according to any one of claims 1 to 5.
- A composition comprising the triazinyl group- or pyrimidinyl group-containing silicone derivative according to any one of claims 1 to 5, and silicone oil and/or liquid oil.
  - A base comprising the triazinyl group- or pyrimidinyl group-containing silicone derivative according to any one of claims 1 to 5.

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